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ION-SENSITIVE, WATER-DISPERSIBLE, CATIONIC POLYMERS, A
METHOD OF MAKING SAME AND ITEMS USING SAME

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FIELD OF THE INVENTION

10 The present invention is directed to ion-sensitive, water-
dispersible cationic polymers and polymer formulations. The present
invention is also directed to a method of making ion-sensitive, water-
dispersible cationic polymers and polymer formulations and their applicability
as binder compositions for disposable items. The present invention is further
15 directed to disposable items, such as wet-wipes comprising ion-sensitive,
water-dispersible binder compositions.

BACKGROUND OF THE INVENTION

20 For many years, the problem of disposability has plagued
industries which provide disposable items, such as, diapers, wet wipes,
incontinent garments and feminine care products. While much headway has
been made in addressing this problem, one of the weak links has been the
inability to create an economical coherent fibrous web, which will readily
dissolve or disintegrate in water, but still have sufficient in-use strength. See,
for example, U.K. patent disclosure 2,241,373 and U.S. Pat. No. 4,186,233.
25 Without such a product, the ability of the user to dispose of the product by
flushing it down the toilet is greatly reduced, if not eliminated. Furthermore,
the ability of the product to disintegrate in a landfill is quite limited because a
large portion of the product components, which may well be biodegradable or
photodegradable, are encapsulated in or bound together by plastic which
30 degrades over a long period of time, if at all. Accordingly, if the plastic

disintegrated in the presence of water, the internal components could degrade as a result of the rupture of the plastic encapsulation or binding.

Disposable products, such as diapers, feminine care products and adult incontinent care products may be made to be disposed by flushing down toilets. Usually such products comprise a body side liner which must rapidly pass fluids, such as urine or menses, so that the fluid may be absorbed by an absorbent core of the product. Typically, the body side liner may be a coherent fibrous web, which desirably possesses a number of characteristics, such as softness and flexibility. The fibrous web of the body side liner material may be typically formed by wet or dry (air) laying a generally random plurality of fibers and joining them together to form a coherent web with a binder compositions. Past binder compositions have preformed this function well. However, fibrous webs comprising these compositions tended to be non-dispersible and present problems in typical household sanitation systems.

Recent binder compositions have been developed which can be more dispersible and are more environmentally responsible than past binder compositions. One class of binder compositions includes polymeric materials having inverse solubility in water. These binder compositions are insoluble in warm water, but are soluble in cold water, such as found in a toilet. It is well known that a number of polymers exhibit cloud points or inverse solubility properties in aqueous media. These polymers have been cited in several publications for various applications, including (1) as evaporation retarders (JP 6207162); (2) as temperature sensitive compositions, which are useful as temperature indicators due to a sharp color change associated with a corresponding temperature change (JP 6192527); (3) as heat sensitive materials that are opaque at a specific temperature and become transparent when cooled to below the specific temperature (JP 51003248 and JP 81035703); (4) as wound dressings with good absorbing characteristics and easy removal (JP 6233809); and (5) as materials in flushable personal care products (U.S. Pat. No. 5,509,913, issued to Richard S. Yeo on April 23, 1996 and assigned to Kimberly-Clark Corporation).

Other recent binders of interest include a class of binders, which are ion-sensitive. Several U.S. and European patents assigned to Lion Corporation of Tokyo, Japan, disclose ion-sensitive polymers comprising acrylic acid and alkyl or aryl acrylates. See U.S. Pat. Nos. 5,312,883, 5,317,063 and 5,384,189, the disclosures of which are incorporated herein by reference, as well as, European Pat. No. 608460A1. In U.S. Pat. No. 5,312,883, terpolymers are disclosed as suitable binders for flushable nonwoven webs. The disclosed acrylic acid-based terpolymers, which comprise partially neutralized acrylic acid, butyl acrylate and 2-ethylhexyl acrylate, are suitable binders for use in flushable nonwoven webs in some parts of the world. However, because of the presence of a small amount of sodium acrylate in the partially neutralized terpolymer, these binders fail to disperse in water containing more than about 15 ppm Ca^{2+} and/or Mg^{2+} . When placed in water containing more than about 15 ppm Ca^{2+} and/or Mg^{2+} ions, nonwoven webs using the above-described binders maintain a tensile strength greater than 30 g/in, which negatively affects the "dispersibility" of the web. The proposed mechanism for the failure is that each calcium ion binds with two carboxylate groups either intramolecularly or intermolecularly. Intramolecular association causes the polymer chain to coil up, which eventually leads to polymer precipitation. Intermolecular association yields crosslinking. Whether intramolecular or intermolecular associations are taking place, the terpolymer is not soluble in water containing more than about 15 ppm Ca^{2+} and/or Mg^{2+} . Due to the strong interaction between calcium ions and the carboxylate groups of the terpolymer, dissociation of the complex is highly unlikely because this association is irreversible. Therefore, the above-described polymer that has been exposed to a high Ca^{2+} and/or Mg^{2+} concentration solution will not disperse in water even if the calcium concentration decreases. This limits the application of the polymer as a flushable binder material because most areas across the U.S. have hard water, which contains more than 15 ppm Ca^{2+} and/or Mg^{2+} .

In a co-pending application assigned to Kimberly Clark; *i.e.*, U.S. patent application Serial No. 09/223,999, filed December 31, 1998, the disclosure of which is incorporated herein by reference, there is disclosed a modification of the acrylic acid terpolymers of the above-referenced patents to
5 Lion Corporation. Specifically, U.S. patent application Serial No. 09/223,999 discloses a sulfonate anion modified acrylic acid terpolymers which has improved dispersibility in relatively hard water; *e.g.*, up to 200 ppm Ca^{2+} and/or Mg^{2+} , compared to the unmodified Lion polymers. However, the Lion Corporation ion-sensitive polymers of the above-referenced patents and the
10 sulfonate anion modified acrylic acid terpolymers of the co-pending application, when used as binders for personal care products, such as wet wipes, typically have reduced sheet wettability, increased sheet stiffness, increased sheet stickiness, reduced binder sprayability and relatively high product cost.

15 Another approach to dispersible personal care products is disclosed in U.S. Pat. No. 5,281,306 to Kao Corporation of Tokyo, Japan. This patent discloses a water-disintegratable cleansing sheet; *i.e.*, wet wipe, comprising water-dispersible fibers treated with a water-soluble binder having a carboxyl group. The cleansing sheet is treated with a cleansing agent
20 containing 5%-95% of a water-compatible organic solvent and 95%-5% water. A preferred organic solvent is propylene glycol. The cleansing sheet retains wet strength and does not disperse in the organic solvent-based cleansing agent, but disperses in water.

25 Although many patents disclose various ion and temperature sensitive compositions for water-dispersible or flushable materials, there exists a need for dispersible products possessing softness, flexibility, three dimensionality, and resiliency; wicking and structural integrity in the presence of body fluids (including feces) at body temperature; and true fiber dispersion after toilet flushing so that fibers do not become entangled with tree roots or at
30 bends in sewer pipes. In addition, the known ion-sensitive polymers, such as those of Lion Corporation and the co-pending application of Kimberly Clark,

have relatively high viscosities at high shear rates that make application by spraying impossible or impractical. Moreover, there is a need in the art for flushable products having water-dispersibility in all areas of the world, including soft and hard water areas. Furthermore, there is a need for water-dispersible binders that do not reduce wettability of product with which they are used and are sprayable for easy and uniform application to and penetration into products. Finally, there is a need for water-dispersible, flushable wet wipes that are stable during storage and retain a desired level of wet strength during use and are wetted with a wetting composition that is relatively free, or is substantially free, of organic solvents. Such a product is needed at a reasonable cost without compromising product safety and environmental concerns, something that past products have failed to do.

SUMMARY OF THE INVENTION

The present invention is directed to ion-sensitive cationic polymers and polymer formulations, which have been developed to address the above-described problems associated with currently available, ion-sensitive polymers and other polymers described in literature. The ion-sensitive polymer formulations of the present invention have a "trigger property," such that the polymers are insoluble in a wetting composition comprising ions of a particular type and concentration, such as monovalent salt solutions at concentrations above about 2%, but are soluble when diluted with water including divalent salt solutions such as hard water with up to 200 ppm (parts per million) calcium and magnesium ions. Unlike some ion-sensitive polymer formulations, which lose dispersibility in hard water because of ion cross-linking by calcium ions, the cationic polymer formulations of the present invention are insensitive to calcium and/or magnesium ions. Consequently, flushable products containing the polymer formulations of the present invention maintain dispersibility in hard water.

The polymer formulations of the present invention are useful as binders and structural components for air-laid and wet-laid nonwoven fabrics

for applications such as body-side liners, fluid distribution materials, fluid intake materials (surge) or cover stock in various personal care products. The polymer formulations of the present invention are particularly useful as a binder material for flushable personal care products, particularly wet wipes for personal use, such as cleaning or treating skin, make-up removal, nail polish removal, medical care, and also wipes for use in hard surface cleaning, automotive care, including wipes comprising cleaning agents, disinfectants, and the like. The flushable products maintain integrity or wet strength during storage and use, and break apart or disperse after disposal in the toilet when the salt concentration falls below a critical level. Suitable substrates for treatment include tissue, such as creped or uncreped tissue, coform products, hydroentangled webs, airlaid mats, fluff pulp, nonwoven webs, and composites thereof. Methods for producing uncreped tissues and molded three-dimensional tissue webs of use in the present invention can be found in commonly owned U.S. patent application, Serial Number 08/912,906, "Wet Resilient Webs and Disposable Articles Made Therewith," by F.-J. Chen et al., filed Aug. 15, 1997; U.S. Pat. No. 5,429,686, issued to Chiu et al. on July 4, 1995; U.S. Pat. No. 5,399,412, issued to S.J. Sudall and S.A. Engel on March 21, 1995; U.S. Pat. No. 5,672,248, issued to Wendt et al. on Sept. 30, 1997; and U.S. Pat. No. 5,607,551, issued to Farrington et al. on March 4, 1997; all of which are incorporated herein by reference in their entirety. The molded tissue structures of the above patents can be especially helpful in providing good cleaning in a wet wipe. Good cleaning can also be promoted by providing a degree of texture in other substrates as well by embossing, molding, wetting and through-air drying on a textured fabric, and the like.

Airlaid material can be formed by metering an airflow containing the fibers and other optional materials, in substantially dry condition, onto a typically horizontally moving wire forming screen. Suitable systems and apparatus for air-laying mixtures of fibers and thermoplastic material are disclosed in, for example, U.S. Pat. No. 4,157,724 (Persson), issued Jun. 12, 1979, and reissued Dec. 25, 1984 as Re. U.S. Pat. No. 31,775;

U.S. Pat. No. 4,278,113 (Persson), issued Jul. 14, 1981; U.S. Pat. No. 4,264,289 (Day), issued Apr. 28, 1981; U.S. Pat. No. 4,352,649 (Jacobsen et al.), issued Oct. 5, 1982; U.S. Pat. No. 4,353,687 (Hosler, et al.), issued Oct. 12, 1982; U.S. Pat. No. 4,494,278 (Kroyer, et al.), issued Jan. 22, 1985; U.S. Pat. No. 4,627,806 (Johnson), issued Dec. 9, 1986; U.S. Pat. No. 4,650,409 (Nistri, et al.), issued Mar. 17, 1987; and U.S. Pat. No. 4,724,980 (Farley), issued Feb. 16, 1988; and U.S. Pat. No. 4,640,810 (Laursen et al.), issued Feb. 3, 1987, the disclosures of which are all incorporated herein by reference.

The present invention also discloses how to make water-dispersible nonwovens, including cover stock (liner), intake (surge) materials and wet wipes, which are stable in fluids having a first ionic composition, such as monovalent ions at a particular concentration substantially greater than is found in typical hard water, using the above-described unique polymer formulations as binder compositions. The resultant nonwovens are flushable and water-dispersible due to the tailored ion sensitivity, which can be triggered regardless of the hardness of water found in toilets throughout the United States and the world.

The present invention further discloses an improved wetting composition for wet wipes. Wet wipes employing the polymer formulations of the present invention are stable during storage and retain a desired level of wet strength during use and are wetted with a wetting composition or cleaning agent that can be relatively free, or is substantially free, of organic solvents.

These features and advantages of the present invention will become apparent after a review of the following detailed description of the disclosed embodiments and the appended drawing and claims.

DETAILED DESCRIPTION OF THE DISCLOSED EMBODIMENTS

In order to be an effective ion-sensitive polymer or polymer formulation suitable for use in flushable or water-dispersible personal care products, the formulations should desirably be (1) functional; *i.e.*, maintain wet strength under controlled conditions and dissolve or disperse rapidly in

soft or hard water, such as found in toilets and sinks around the world; (2) safe (not toxic); and (3) relatively economical. In addition to the foregoing factors, the ion-sensitive formulations when used as a binder composition for a non-woven substrate, such as a wet wipe, desirably should be (4) processable on a commercial basis; *i.e.*, may be applied relatively quickly on a large scale basis, such as by spraying, which thereby requires that the binder composition have a relatively low viscosity at high shear; and (5) provide acceptable levels of sheet or substrate wettability. The wetting composition with which the wet wipes of the present invention are treated can provide some of the foregoing advantages, and, in addition, can provide one or more of (6) improved skin care, such as reduced skin irritation or other benefits, (7) improved tactile properties, and (8) promote good cleaning by providing a balance in use between friction and lubricity on the skin (skin glide). The ion-sensitive cationic polymers and polymer formulations of the present invention and articles made therewith, especially wet wipes comprising particular wetting compositions set forth below, can meet many or all of the above criteria. Of course, it is not necessary for all of the advantages of the preferred embodiments of the present invention to be met to fall within the scope of the present invention.

The cationic polymers of the present invention may be formed from four monomers different monomers. The quaternary polymers of the present invention are the polymerization product of the following four monomers: acrylic acid, butyl acrylate, 2-ethylhexyl acrylate and [2-(methacryloyloxy)ethyl] trimethyl ammonium chloride. In an alternate embodiment, the cationic polymers are formed from four different monomers. The quaternary polymers of the present invention are the polymerization product of the following four monomers: acrylamide, butyl acrylate, 2-ethylhexyl acrylate and [2-(methacryloyloxy)ethyl] trimethyl ammonium chloride. In yet another embodiment of the present invention, the cationic polymers are formed from three different polymers. The terpolymers of the present invention are the polymerization product of the following three

monomers: butyl acrylate, 2-ethylhexyl acrylate and [2-(methacryloyloxy)ethyl] trimethyl ammonium chloride.

Ion-sensitive cationic polymers of the present invention may
5 comprise any combination of the above-referenced monomers capable of free
radical polymerization into a terpolymer or quaternary polymer. Suitable
acrylic acid monomers include, but are not limited to, acrylic acid and
methacrylic acid. Suitable acrylic monomers include, but are not limited to,
acrylic esters and methacrylic esters having an alkyl group of 1 to 18 carbon
10 atoms or a cycloalkyl group of 3 to 18 carbon atoms and it is preferred that
acrylic esters and/or methacrylic esters having a alkyl group of 1 to 12 carbon
atoms or a cycloalkyl group of 3 to 12 carbon atoms be used singly or in
combination. Other suitable monomers include, but are not limited to,
acrylamide and methacrylamide based monomers, such as acrylamide, N,N-
15 dimethyl acrylamide, N-ethyl acrylamide, N-isopropyl acrylamide, and
hydroxymethyl acrylamide; N-vinylpyrrolidinone; N-vinylforamide;
hydroxyalkyl acrylates and hydroxyalkyl methacrylates, such as hydroxyethyl
methacrylate and hydroxyethyl acrylate. Other suitable acrylic acid monomers
are disclosed in U.S. Patent No. 5,317,063, assigned to Lion Corporation,
20 Tokyo, Japan, the disclosure of which is incorporated herein by reference in its
entirety. ¹

The relative amounts of the monomers in the terpolymers and
quaternary polymers of the present invention may vary depending on the
desired properties in the resulting polymer. For the quaternary polymer made
25 from acrylic acid, butyl acrylate, 2-ethylhexyl acrylate and [2-
(methacryloyloxy)ethyl] trimethyl ammonium chloride, the mole percent of
monomer in the quaternary polymer is as follows: __ to __ mole percent
acrylic acid; __ to __ mole percent butyl acrylate; __ to __ mole percent 2-
ethylhexyl acrylate; and __ to __ mole percent [2-(methacryloyloxy)ethyl]
30 trimethyl ammonium chloride. More specifically, the mole percent of

monomers in the quaternary polymer is from about ___ to about ___ mole percent _____. Most specifically, the mole percent of monomers in the quaternary polymer is from about ___ to about ___ mole percent _____.

5 For the quaternary polymer made from acrylamide, butyl acrylate, 2-ethylhexyl acrylate and [2-(methacryloyloxy)ethyl] trimethyl ammonium chloride, the mole percent of monomer in the quaternary polymer is as follows: ___ to ___ mole percent acrylamide; ___ to ___ mole percent butyl acrylate; ___ to ___ mole percent 2-ethylhexyl acrylate; and ___ to ___ mole percent [2-(methacryloyloxy)ethyl] trimethyl ammonium chloride. More
10 specifically, the mole percent of monomers in the quaternary polymer is about ___ to about ___ mole percent _____. Most specifically, the mole percent of monomers in the quaternary polymer is about ___ to about ___ mole percent _____.

15 For the terpolymer made from butyl acrylate, 2-ethylhexyl acrylate and [2-(methacryloyloxy)ethyl] trimethyl ammonium chloride, the mole percent of monomer in the quaternary polymer is as follows: ___ to ___ mole percent butyl acrylate; ___ to ___ mole percent 2-ethylhexyl acrylate; and ___ to ___ mole percent [2-(methacryloyloxy)ethyl] trimethyl ammonium
20 chloride. More specifically, the mole percent of monomers in the terpolymer is about ___ to about ___ mole percent _____. Most specifically, the mole percent of monomers in the terpolymer is about ___ to about ___ mole percent _____.

25 The terpolymers and quaternary polymers of the present invention may have an average molecular weight, which varies depending on the ultimate use of the polymer. The quaternary polymers of the present invention have a weight average molecular weight ranging from about 10,000 to about 5,000,000. More specifically, the quaternary polymers of the present
30 invention have a weight average molecular weight ranging from about 25,000

¹ I need to know commercial sources and brand names for acrylic acid, butyl acrylate, 2-ethylhexyl

to about 2,000,000, or, more specifically still, from about 200,000 to about 1,000,000. The terpolymers of the present invention have a weight average molecular weight ranging from about 10,000 to about 5,000,000. More specifically, the terpolymers of the present invention have a weight average molecular weight ranging from about 25,000 to about 2,000,000, or, more specifically still, from about 200,000 to about 1,000,000.

The terpolymers and quaternary polymers of the present invention may be prepared according to a variety of polymerization methods, desirably a solution polymerization method. Suitable solvents for the polymerization method include, but are not limited to, lower alcohols such as methanol, ethanol and propanol; a mixed solvent of water and one or more lower alcohols mentioned above; and a mixed solvent of water and one or more lower ketones such as acetone or methyl ethyl ketone.

In the polymerization methods of the present invention, any free radical polymerization initiator may be used. Selection of a particular initiator may depend on a number of factors including, but not limited to, the polymerization temperature, the solvent, and the monomers used. Suitable polymerization initiators for use in the present invention include, but are not limited to, 2,2'-azobisisobutyronitrile, 2,2'-azobis(2-methylbutyronitrile), 2,2'-azobis(2,4-dimethylvaleronitrile), 2,2'-azobis(2-amidinopropane)dihydrochloride, 2,2'-azobis(N,N'-dimethyleneisobutylamidine), potassium persulfate, ammonium persulfate, and aqueous hydrogen peroxide. The amount of polymerization initiator may desirably range from about 0.01 to 5 weight percent based on the total weight of monomer present.

The polymerization temperature may vary depending on the polymerization solvent, monomers, and initiator used, but in general, ranges from about 20 °C. to about 90 °C. Polymerization time generally ranges from about 2 to about 8 hours.

The cationic polymers in accordance with the present invention include hydrophilic monomers, such as acrylic acid or methacrylic acid, incorporated into the polymers of the present invention along with _____ . The sulfonate anions of these monomers are softer than carboxylate anions since the negative charge of the sulfonate anion is delocalized over three oxygen atoms and a larger sulfur atom, as opposed to only two oxygen atoms and a smaller carbon atom in the carboxylate anion. These monomers, containing the softer sulfonate anion, are less interactive with multivalent ions present in hard water, particularly Ca^{2+} and Mg^{2+} ions. Suitable _____ To maintain the hydrophobic/hydrophilic balance of the ion-sensitive polymer, one or more hydrophobic monomers are added to the polymer.

In a further embodiment of the present invention, the above-described ion-sensitive polymer formulations are used as binder materials for flushable and/or non-flushable products. In order to be effective as a binder material in flushable products throughout the United States, the ion-sensitive polymer formulations of the present invention remain stable and maintain their integrity while dry or in relatively high concentrations of monovalent ions, but become soluble in water containing up to about 200 ppm or more divalent ions, especially calcium, magnesium and sulfonates ions. Desirably, the ion-sensitive polymer formulations of the present invention are insoluble in a salt solution containing at least about 2 weight percent of one or more inorganic and/or organic salts containing monovalent ions. More desirably, the ion-sensitive polymer formulations of the present invention are insoluble in a salt solution containing from about 2 weight percent to about ____ weight percent of one or more inorganic and/or organic salts containing monovalent ions. Even more desirably, the ion-sensitive polymer formulations of the present invention are insoluble in salt solutions containing from about 2 weight percent to about ____ weight percent of one or more inorganic and/or organic salts containing monovalent ions. Suitable monovalent ions include, but are not limited to, Na^+ ions, K^+ ions, Li^+ ions, NH_4^+ ions, Zn^{2+} ions, low

molecular weight quaternary ammonium compounds (e.g., those having fewer than 5 carbons on any side group), and a combination thereof.

Based on a recent study conducted by the American Chemical Society, water hardness across the United States varies greatly, with CaCO_3 concentration ranging from near zero for soft water to about 500 ppm CaCO_3 (about 200 ppm Ca^{2+} ion) for very hard water. To ensure polymer formulation dispersibility across the country (and throughout the whole world), the ion-sensitive polymer formulations of the present invention are desirably soluble in water containing up to about 50 ppm Ca^{2+} and/or Mg^{2+} ions. More desirably, the ion-sensitive polymer formulations of the present invention are soluble in water containing up to about 100 ppm Ca^{2+} and/or Mg^{2+} ions. Even more desirably, the ion-sensitive polymer formulations of the present invention are soluble in water containing up to about 150 ppm Ca^{2+} and/or Mg^{2+} ions. Even more desirably, the ion-sensitive polymer formulations of the present invention are soluble in water containing up to about 200 ppm Ca^{2+} and/or Mg^{2+} ions.

The co-binder polymer can comprise surface active compounds that improve the wettability of the substrate after application of the binder mixture. Wettability of a dry substrate that has been treated with an ion-sensitive polymer formulation can be a problem in some embodiments, because the hydrophobic portions of the ion-sensitive polymer formulation can become selectively oriented toward the air phase during drying, creating a hydrophobic surface that can be difficult to wet when the wetting composition is later applied unless surfactants are added to the wetting composition. Surfactants, or other surface active ingredients, in polymers can improve the wettability of the dried substrate that has been treated with an ion-sensitive polymer formulation. Surfactants in the polymer should not significantly interfere with the ion-sensitive polymer formulation. Thus, the binder should maintain good integrity and tactile properties in the pre-moistened wipes with the surfactant present.

Binder Formulations and Fabrics Containing the Same

The polymer formulations of the present invention may be used as binders. The binder formulations of the present invention may be applied to any fibrous substrate. The binders are particularly suitable for use in water-dispersible products. Suitable fibrous substrates include, but are not limited to, nonwoven and woven fabrics. In many embodiments, particularly personal care products, preferred substrates are nonwoven fabrics. As used herein, the term "nonwoven fabric" refers to a fabric that has a structure of individual fibers or filaments randomly arranged in a mat-like fashion (including papers). Nonwoven fabrics can be made from a variety of processes including, but not limited to, air-laid processes, wet-laid processes, hydroentangling processes, staple fiber carding and bonding, and solution spinning.

The binder composition may be applied to the fibrous substrate by any known process of application. Suitable processes for applying the binder material include, but are not limited to, printing, spraying, electrostatic spraying, coating, flooded nips, metered press rolls, impregnating or by any other technique. The amount of binder composition may be metered and distributed uniformly within the fibrous substrate or may be non-uniformly distributed within the fibrous substrate. The binder composition may be distributed throughout the entire fibrous substrate or it may be distributed within a multiplicity of small closely spaced areas. In most embodiments, uniform distribution of binder composition is desired.

For ease of application to the fibrous substrate, the binder may be dissolved in water, or in a non-aqueous solvent, such as methanol, ethanol, acetone, or the like, with water being the preferred solvent. The amount of binder dissolved in the solvent may vary depending on the polymer used and the fabric application. Desirably, the binder solution contains up to about 25 percent by weight of binder composition solids. More desirably, the binder solution contains from about 10 to 20 percent by weight of binder composition solids, especially about 12 percent by weight binder composition solids. Plasticizers, perfumes, coloring agents, antifoams, bactericides, preservative,

surface active agents, thickening agents, fillers, opacifiers, tackifiers, detackifiers, and similar additives can be incorporated into the solution of binder components, if so desired.

5 Once the binder composition is applied to the substrate, the substrate is dried by any conventional means. Once dry, the coherent fibrous substrate exhibits improved tensile strength when compared to the tensile strength of the untreated wet-laid or dry-laid substrates, and yet has the ability to rapidly "fall apart", or disintegrate when placed in soft or hard water having a relatively high multivalent ionic concentration and agitated. For example,
10 the dry tensile strength of the fibrous substrate may be increased by at least 25 percent as compared to the dry tensile strength of the untreated substrate not containing the binder. More particularly, the dry tensile strength of the fibrous substrate may be increase by at least 100 percent as compared to the dry tensile strength of the untreated substrate not containing the binder. Even
15 more particularly, the dry tensile strength of the fibrous substrate may be increased by at least 500 percent as compared to the dry tensile strength of the untreated substrate not containing the binder.

 A desirable feature of the present invention is that the improvement in tensile strength is effected where the amount of binder composition present, "add-on", in the resultant fibrous substrate represents
20 only a small portion by weight of the entire substrate. The amount of "add-on" can vary for a particular application; however, the optimum amount of "add-on" results in a fibrous substrate which has integrity while in use and also quickly disperses when agitated in water. For example, the binder components typically are from about 5 to about 65 percent, by weight, of the
25 total weight of the substrate. More particularly, the binder components may be from about 10 to about 35 percent, by weight, of the total weight of the substrate. Even more particularly, the binder components may be from about 17 to about 22 percent by weight of the total weight of the substrate.

30 The nonwoven fabrics of the present invention have good in-use tensile strength, as well as, ion triggerability. Desirably, the nonwoven

fabrics of the present invention are abrasion resistant and retain significant tensile strength in aqueous solutions containing greater than about 2 weight percent monovalent ions, such as NaCl, KCl, LiCl, a mixture of monovalent ions or a mixture of monovalent and multivalent ions, such as NaCl and ZnCl₂. Because of this latter property, nonwoven fabrics of the present invention are well suited for disposable products, such as sanitary napkins, diapers, adult incontinence products, and dry and premoistened wipes (wet wipes), which can be thrown in a flush toilet after use in any part of the world.

The fibers forming the fabrics above can be made from a variety of materials including natural fibers, synthetic fibers, and combinations thereof. The choice of fibers depends upon, for example, the intended end use of the finished fabric and fiber cost. For instance, suitable fibrous substrates may include, but are not limited to, natural fibers such as cotton, linen, jute, hemp, wool, wood pulp, etc. Similarly, regenerated cellulosic fibers, such as viscose rayon and cuprammonium rayon, modified cellulosic fibers, such as cellulose acetate, or synthetic fibers, such as those derived from polypropylenes, polyethylenes, polyolefins, polyesters, polyamides, polyacrylics, etc., alone or in combination with one another, may likewise be used. Blends of one or more of the above fibers may also be used, if so desired. Among wood pulp fibers, any known papermaking fibers may be used, including softwood and hardwood fibers. Fibers, for example, may be chemically pulped or mechanically pulped, bleached or unbleached, virgin or recycled, high yield or low yield, and the like. Mercerized, chemically stiffened or crosslinked fibers may also be used.

Synthetic cellulose fiber types include rayon in all its varieties and other fibers derived from viscose or chemically modified cellulose, including regenerated cellulose and solvent-spun cellulose, such as Lyocell. Chemically treated natural cellulosic fibers can be used, such as mercerized pulps, chemically stiffened or crosslinked fibers, or sulfonated fibers. Recycled fibers, as well as virgin fibers, can be used. Cellulose produced by microbes and other cellulosic derivatives can be used. As used herein, the

term "cellulosic" is meant to include any material having cellulose as a major constituent, and, specifically, comprising at least 50 percent by weight cellulose or a cellulose derivative. Thus, the term includes cotton, typical wood pulps, non-woody cellulosic fibers, cellulose acetate, cellulose triacetate, rayon, thermomechanical wood pulp, chemical wood pulp, debonded chemical wood pulp, milkweed, or bacterial cellulose. The fiber length is important in producing the fabrics of the present invention. In some embodiments, such as flushable products, fiber length is of more importance. The minimum length of the fibers depends on the method selected for forming the fibrous substrate. For example, where the fibrous substrate is formed by carding, the length of the fiber should usually be at least about 42 mm in order to insure uniformity. Where the fibrous substrate is formed by air-laid or wet-laid processes, the fiber length may desirably be about 0.2 to 6 mm. Although fibers having a length of greater than 50 mm are within the scope of the present invention, it has been determined that when a substantial quantity of fibers having a length greater than about 15 mm is placed in a flushable fabric, though the fibers will disperse and separate in water, their length tends to form "ropes" of fibers, which are undesirable when flushing in home toilets. Therefore, for these products, it is desired that the fiber length be about 15 mm or less so that the fibers will not have a tendency to "rope" when they are flushed through a toilet. Although fibers of various lengths are applicable in the present invention, desirably fibers are of a length less than about 15 mm so that the fibers disperse easily from one another when in contact with water. The fibers, particularly synthetic fibers, can also be crimped

The fabrics of the present invention may be formed from a single layer or multiple layers. In the case of multiple layers, the layers are generally positioned in a juxtaposed or surface-to-surface relationship and all or a portion of the layers may be bound to adjacent layers. Nonwoven webs of the present invention may also be formed from a plurality of separate nonwoven webs wherein the separate nonwoven webs may be formed from single or multiple layers. In those instances where the nonwoven web

includes multiple layers, the entire thickness of the nonwoven web may be subjected to a binder application or each individual layer may be separately subjected to a binder application and then combined with other layers in a juxtaposed relationship to form the finished nonwoven web.

5 In one embodiment, the fabric substrates of the present invention may be incorporated into cleansing and body fluid absorbent products, such as sanitary napkins, diapers, adult incontinence products, surgical dressings, tissues, wet wipes, and the like. These products may include an absorbent core, comprising one or more layers of an absorbent
10 fibrous material. The core may also comprise one or more layers of a fluid-pervious element, such as fibrous tissue, gauze, plastic netting, etc. These are generally useful as wrapping materials to hold the components of the core together. Additionally, the core may comprise a fluid-impervious element or barrier means to preclude the passage of fluid through the core and on the
15 outer surfaces of the product. Desirably, the barrier means also is water-dispersible. A film of a polymer having substantially the same composition as the aforesaid water-dispersible binder is particularly well-suited for this purpose. In accordance with the present invention, the polymer compositions are useful for forming each of the above-mentioned product components
20 including the layers of absorbent core, the fluid-pervious element, the wrapping materials, and the fluid-impervious element or barrier means.

 The binder formulations of the present invention are particularly useful for binding fibers of air-laid nonwoven fabrics. These air-laid materials are useful for body-side liners, fluid distribution materials, fluid
25 in-take materials, such as a surge material, absorbent wrap sheet and cover stock for various water-dispersible personal care products. Air-laid materials are particularly useful for use as a pre-moistened wipe (wet wipe). The basis weights for air-laid non-woven fabrics may range from about 20 to about 200 grams per square meter ("gsm") with staple fibers having a denier of about
30 0.5-10 and a length of about 6-15 millimeters. Surge, or in-take, materials need better resiliency and higher loft so staple fibers having about 6 denier or

greater are used to make these products. A desirable final density for the surge, or in-take, materials is between about 0.025 grams per cubic centimeter ("g/cc") to about 0.10 g/cc. Fluid distribution materials may have a higher density, in the desired range of about 0.10 to about 0.20 g/cc using fibers of lower denier, most desirable fibers have a denier of less than about 1.5. Wipes generally can have a fiber density of about 0.025 g/cc to about 0.2 g/cc and a basis weight of about 20 gsm to about 150 gsm; specifically from about 30 to about 90 gsm, and most specifically from about 60 gsm to about 65 gsm.

The nonwoven fabrics of the present invention may also be incorporated into such body fluid absorbing products as sanitary napkins, diapers, surgical dressings, tissues and the like. In one embodiment, the binder is such that it will not dissolve when contacted by body fluids since the concentration of monovalent ions in the body fluids is above the level needed for dissolution; *i.e.*, greater than 2% by weight. The nonwoven fabric retains its structure, softness and exhibits a toughness satisfactory for practical use. However, when brought into contact with water having a concentration of multivalent ions, such as Ca^{2+} and Mg^{2+} ions, of up to about 200 ppm or more, the binder disperses. The nonwoven fabric structure is then easily broken and dispersed in the water.

In one embodiment of the present invention, the in-use tensile strength of a nonwoven fabric is enhanced by forming the nonwoven fabric with a binder material comprising the ion-sensitive polymer formulation of the present invention and subsequently applying one or more monovalent and/or multivalent salts to the nonwoven fabric. The salt may be applied to the nonwoven fabric by any method known to those of ordinary skill in the art including, but not limited to, applying a solid powder onto the fabric and spraying a salt solution onto the fabric. The amount of salt may vary depending on a particular application. However, the amount of salt applied to the fabric is typically from about 2 weight percent to about 10 weight percent salt solids based on the total weight of the fabric. The salt-containing fabrics

of the present invention may be used in a variety of fabric applications including, but not limited to, feminine pads, surgical dressings, and diapers.

Those skilled in the art will readily understand that the binder formulations and fibrous substrates of the present invention may be advantageously employed in the preparation of a wide variety of products, including but not limited to, absorbent personal care products designed to be contacted with body fluids. Such products may only comprise a single layer of the fibrous substrate, or may comprise a combination of elements, as described above. Although the binder formulations and fibrous substrates of the present invention are particularly suited for personal care products, the binder formulations and fibrous substrates may be advantageously employed in a wide variety of consumer products.

Unlike other binder systems known in the art, the ion-sensitive polymer formulations of the present invention can be activated as binders without the need for elevated temperature. While drying or water removal is useful in achieving a good distribution of the binder in a fibrous web, elevated temperature, per se, is not essential because the binder does not require crosslinking or other chemical reactions with high activation energy to serve as a binder. Rather, the interaction with a soluble activating compound, typically a salt, is sufficient to cause the binder to become active (insoluble) or "salted out." Thus, a drying step can be avoided, if desired, or replaced with low-temperature water removal operations such as room-temperature drying or freeze drying. Elevated temperature is generally helpful for drying, but the drying can be done at temperatures below what is normally needed to drive crosslinking reactions. Thus, the peak temperature to which the substrate is exposed or to which the substrate is brought can be below any of the following: 180° C., 160° C., 140° C., 120° C., 110° C., 105° C., 100° C., 90° C., 75° C., and 60° C., with an exemplary range for peak web temperature of from about 50° C. to about 110° C., or from about 70° C. to about 140° C. Of course, higher temperatures can be used, but are not necessary in most embodiments. While polymer systems, such as commercial latex emulsions,

may also comprise crosslinkers suited for reaction at temperatures of 160° C. or higher, maintaining a lower peak temperature can be beneficial in preventing development of excessive strength in the polymer that might otherwise hinder the water dispersibility of the pre-moistened wipe.

5

Wet Wipe Wetting Composition and Wet Wipes Containing the Same

One particularly interesting embodiment of the present invention is the production of pre-moistened wipes, or wet wipes, from the above-described ion-sensitive binder compositions and fibrous materials. For
10 wipes, the fibrous material may be in the form of a woven or nonwoven fabric; however, nonwoven fabrics are more desirable. The nonwoven fabric is desirably formed from relatively short fibers, such as wood pulp fibers. The minimum length of the fibers depends on the method selected for forming the nonwoven fabric. Where the nonwoven fabric is formed by a wet or dry
15 method, the fiber length is desirably from about 0.1 millimeters to 15 millimeters. Desirably, the nonwoven fabric of the present invention has a relatively low wet cohesive strength when it is not bonded together by an adhesive or binder material. When such nonwoven fabrics are bonded together by a binder composition, which loses its bonding strength in tap water
20 and in sewer water, the fabric will break up readily by the agitation provided by flushing and moving through the sewer pipes.

The finished wipes may be individually packaged, desirably in a folded condition, in a moisture proof envelope or packaged in containers holding any desired number of sheets in a water-tight package with a wetting
25 composition applied to the wipe. The finished wipes may also be packaged as a roll of separable sheets in a moisture-proof container holding any desired number of sheets on the roll with a wetting composition applied to the wipes. The roll can be coreless and either hollow or solid. Coreless rolls, including rolls with a hollow center or without a solid center, can be produced with
30 known coreless roll winders, including those of SRP Industry, Inc. (San Jose, CA); Shimizu Manufacturing (Japan), and the devices disclosed in U.S. Pat.

No. 4,667,890, issued May 26, 1987 to Gietman. Solid-wound coreless rolls can offer more product for a given volume and can be adapted for a wide variety of dispensers.

5 Relative to the weight of the dry fabric, the wipe may desirably contain from about 10 percent to about 400 percent of the wetting composition, more desirably from about 100 percent to about 300 percent of the wetting composition, and even more desirably from about 180 percent to about 240 percent of the wetting composition. The wipe maintains its desired characteristics over the time periods involved in warehousing, transportation,
10 retail display and storage by the consumer. Accordingly, shelf life may range from two months to two years.

 Various forms of impermeable envelopes and storage means for containing wet-packaged materials such as wipes and towelettes and the like are well known in the art. Any of these may be employed in packaging
15 the pre-moistened wipes of the present invention.

 Desirably, the pre-moistened wipes of the present invention are wetted with an aqueous wetting composition, which has one or more of the following properties:

 (1) is compatible with the above-described ion-sensitive
20 binder compositions of the present invention;

 (2) enables the pre-moistened wipe to maintain its wet strength during converting, storage and usage (including dispensing), as well as, dispersibility in a toilet bowl;

 (3) does not cause skin irritation;

25 (4) reduces tackiness of the wipe, and provides unique tactile properties, such as skin glide and a "lotion-like feel"; and

 (5) acts as a vehicle to deliver "moist cleansing" and other skin health benefits.

 The wetting composition should not act as a solvent for the
30 binder and generally does not contain solvents other than water, and particularly does not contain organic solvents, though a small quantity (<1%)

of a fragrance solubilizer, such as polysorbate 20, may be present, depending on the fragrance and the salt concentration of the wetting composition. Desirably, the wetting composition contains less than about 10 weight percent of organic solvents, such as propylene glycol or other glycols, polyhydroxy alcohols, and the like, based on the total weight of the wetting composition. More desirably, the wetting composition contains less than about 4 weight percent of organic solvents. Even more desirably, the wetting composition contains less than about 1 weight percent of organic solvents. The wetting composition can be substantially free of organic solvents.

One aspect of the present invention is a wetting composition, which contains an activating compound that maintains the strength of a water-dispersible binder until the activating compound is diluted with water, whereupon the strength of the water-dispersible binder begins to decay. The water-dispersible binder may be any of the ion-sensitive binder compositions of the present invention or any other ion-sensitive binder composition. The activating compound in the wetting composition can be a salt, such as sodium chloride, or any other compound, which provides in-use and storage strength to the water-dispersible binder composition, and can be diluted in water to permit dispersion of the substrate as the binder polymer triggers to a weaker state. Desirably, the wetting composition contains more than about 2 weight percent of an activating compound based on the total weight of the wetting composition. Specifically, the wetting composition may contain from about 2 weight percent to about 20 weight percent of an activating compound. Even more specifically, the wetting composition may contain from about 2 weight percent to about 4 weight percent of an activating compound.

The wetting composition of the present invention may further comprise a variety of additives compatible with the activating compound and the water-dispersible binder, such that the strength and dispersibility functions of the wipe are not jeopardized. Suitable additives in the wetting composition include, but are not limited to, the following additives: skin-care additives; odor control agents; detackifying agents to reduce the tackiness of the binder;

particulates; antimicrobial agents; preservatives; wetting agents and cleaning agents such as detergents, surfactants, and some silicones; emollients; surface feel modifiers for improved tactile sensation (e.g., lubricity) on the skin; fragrance; fragrance solubilizers; opacifiers; fluorescent whitening agents; UV absorbers; pharmaceuticals; and pH control agents, such as malic acid or potassium hydroxide.

Skin-Care Additives

As used herein, the term "skin-care additives" represents additives, which provide one or more benefits to the user, such as a reduction in the probability of having diaper rash and/or other skin damage caused by fecal enzymes. These enzymes, particularly trypsin, chymotrypsin and elastase, are proteolytic enzymes produced in the gastrointestinal tract to digest food. In infants, for example, the feces tend to be watery and contain, among other materials, bacteria, and some amounts of undegraded digestive enzymes. These enzymes, if they remain in contact with the skin for any appreciable period of time, have been found to cause an irritation that is uncomfortable in itself and can predispose the skin to infection by microorganisms. As a countermeasure, skin-care additives include, but are not limited to, the enzyme inhibitors and sequestrants set forth hereafter. The wetting composition may contain less than about 5 weight percent of skin-care additives based on the total weight of the wetting composition. More specifically, the wetting composition may contain from about 0.01 weight percent to about 2 weight percent of skin-care additives. Even more specifically, the wetting composition may contain from about 0.01 weight percent to about 0.05 weight percent of skin-care additives.

A variety of skin-care additives may be added to the wetting composition and the pre-moistened wipes of the present invention or included therein. In one embodiment of the present invention, skin-care additives in the form of particles are added to serve as fecal enzyme inhibitors, offering potential benefits in the reduction of diaper rash and skin damage caused by

fecal enzymes. U.S. Pat. No. 6,051,749, issued April 18, 2000 to Schulz et al., the entirety of which is herein incorporated by reference, discloses organophilic clays in a woven or nonwoven web, said to be useful for inhibiting fecal enzymes. Such materials may be used in the present invention, including reaction products of a long chain organic quaternary ammonium compound with one or more of the following clays: montmorillonite, bentonite, beidellite, hectorite, saponite, and stevensite.

Other known enzyme inhibitors and sequestrants may be used as skin-care additives in the wetting composition of the present invention, including those that inhibit trypsin and other digestive or fecal enzymes, and inhibitors for urease. For example, enzyme inhibitors and anti-microbial agents may be used to prevent the formation of odors in body fluids. For example, urease inhibitors, which are also said to play a role in odor absorption, are disclosed by T. Trinh in World Patent Application No. 98/26808, "Absorbent Articles with Odor Control System," published June 25, 1998, the entirety of which is herein incorporated by reference. Such inhibitors may be incorporated into the wetting composition and the pre-moistened wipes of the present invention and include transition metal ions and their soluble salts, such as silver, copper, zinc, ferric, and aluminum salts. The anion may also provide urease inhibition, such as borate, phytate, etc. Compounds of potential value include, but are not limited to, silver chlorate, silver nitrate, mercury acetate, mercury chloride, mercury nitrate, copper metaborate, copper bromate, copper bromide, copper chloride, copper dichromate, copper nitrate, copper salicylate, copper sulfate, zinc acetate, zinc borate, zinc phytate, zinc bromate, zinc bromide, zinc chlorate, zinc chloride, zinc sulfate, cadmium acetate, cadmium borate, cadmium bromide, cadmium chlorate, cadmium chloride, cadmium formate, cadmium iodate, cadmium iodide, cadmium permanganate, cadmium nitrate, cadmium sulfate, and gold chloride.

Other salts that have been disclosed as having urease inhibition properties include ferric and aluminum salts, especially the nitrates, and

bismuth salts. Other urease inhibitors are disclosed by Trinh, including hydroxamic acid and its derivatives; thiourea; hydroxylamine; salts of phytic acid; extracts of plants of various species, including various tannins, e.g. carob tannin, and their derivatives such as chlorogenic acid derivatives; naturally occurring acids such as ascorbic acid, citric acid, and their salts; phenyl phosphoro diamidate/diamino phosphoric acid phenyl ester; metal aryl phosphoramidate complexes, including substituted phosphorodiamidate compounds; phosphoramidates without substitution on the nitrogen; boric acid and/or its salts, including especially, borax, and/or organic boron acid compounds; the compounds disclosed in European Patent Application 408,199; sodium, copper, manganese, and/or zinc dithiocarbamate; quinones; phenols; thiurams; substituted rhodanine acetic acids; alkylated benzoquinones; formarnidine disulphide; 1:3-diketones maleic anhydride; succinamide; phthalic anhydride; pehenic acid; /N,N-dihalo-2-imidazolidinones; N-halo2-oxazolidinones; thio- and/or acyl-phosphoryltnamide and/or substituted derivatives thereof-, thiopyridine-N-oxides, thiopyridines, and thiopyrimidines; oxidized sulfur derivatives of diaminophosphinyl compounds; cyclotriphosphazatriene derivatives; ortho-diaminophosphinyl derivatives of oximes; bromo-nitro compounds; S-aryl and/or alkyl diamidophosphorothiolates; diaminophosphinyl derivatives; mono- and/or polyphosphorodiamide; 5-substituted-benzoxathiol-2-ones; N(diaminophosphinyl)arylcarboxamides; alkoxy-1,2-benzothiazin compounds; etc.

Many other skin-care additives may be incorporated into the wetting composition and pre-moistened wipes of the present invention, including, but not limited to, sun blocking agents and UV absorbers, acne treatments, pharmaceuticals, baking soda (including encapsulated forms thereof), vitamins and their derivatives such as Vitamins A or E, botanicals such as witch hazel extract and aloe vera, allantoin, emollients, disinfectants, hydroxy acids for wrinkle control or anti-aging effects, sunscreens, tanning promoters, skin lighteners, deodorants and anti-perspirants, ceramides for skin

benefits and other uses, astringents, moisturizers, nail polish removers, insect repellants, antioxidants, antiseptics, anti-inflammatory agents and the like, provided that the additives are compatible with an ion-sensitive binder composition associated therewith, and especially the ion-sensitive binder compositions of the present invention (*i.e.*, they do not cause a substantial loss of strength in the wet state of the pre-moistened wipes, prior to dilution in water, while permitting dispersibility in water).

Useful materials for skin care and other benefits are listed in *McCutcheon's 1999*, Vol. 2: Functional Materials, MC Publishing Company, Glen Rock, NJ. Many useful botanicals for skin care are provided by Active Organics, Lewisville, Texas.

Odor Control Additives

Suitable odor control additives for use in the wetting composition and pre-moistened wipes of the present invention include, but are not limited to, zinc salts; talc powder; encapsulated perfumes (including microcapsules, macrocapsules, and perfume encapsulated in liposomes, vessicles, or microemulsions); chelants, such as ethylenediamine tetra-acetic acid; zeolites; activated silica, activated carbon granules or fibers; activated silica particulates; polycarboxylic acids, such as citric acid; cyclodextrins and cyclodextrin derivatives; chitosan or chitin and derivatives thereof; oxidizing agents; antimicrobial agents, including silver-loaded zeolites (*e.g.*, those of BF Technologies, located in Beverly, Massachusetts, sold under the trademark HEALTHSHIELDTM); triclosan; kieselguhr; and mixtures thereof. In addition to controlling odor from the body or body wastes, odor control strategies can also be employed to mask or control any odor of the treated substrate. Desirably, the wetting composition contains less than about 5 weight percent of odor control additives based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 2 weight percent of odor control additives. Even more

desirably, the wetting composition contains from about 0.03 weight percent to about 1 weight percent of odor control additives.

In one embodiment of the present invention, the wetting composition and/or pre-moistened wipes comprise derivatized cyclodextrins, such as hydroxypropyl beta-cyclodextrin in solution, which remain on the skin after wiping and provide an odor-absorbing layer. In other embodiments, the odor source is removed or neutralized by application of an odor-control additive, exemplified by the action of a chelant that binds metal groups necessary for the function of many proteases and other enzymes that commonly produce an odor. Chelating the metal group interferes with the enzyme's action and decreases the risk of malodor in the product.

Principles for the application of chitosan or chitin derivatives to nonwoven webs and cellulosic fibers are described by S. Lee et al. in "Antimicrobial and Blood Repellent Finishes for Cotton and Nonwoven Fabrics Based on Chitosan and Fluoropolymers," Textile Research Journal, 69(2); 104-112, Feb. 1999.

Detackifying Agents

While elevated salt concentrations may reduce the tack of the ion-sensitive binder, other means of tack reduction are often desirable. Thus, detackifying agents may be used in the wetting composition to reduce the tackiness, if any, of the ion-sensitive binder. Suitable detackifiers include any substance known in the art to reduce tack between two adjacent fibrous sheets treated with an adhesive-like polymer or any substance capable of reducing the tacky feel of an adhesive-like polymer on the skin. Detackifiers may be applied as solid particles in dry form, as a suspension or as a slurry of particles. Deposition may be by spray, coating, electrostatic deposition, impingement, filtration (i.e., a pressure differential drives a particle-laden gas phase through the substrate, depositing particles by a filtration mechanism), and the like, and may be applied uniformly on one or more surfaces of the substrate or may be applied in a pattern (e.g., repeating or random patterns)

over a portion of the surface or surfaces of the substrate. The detackifier may be present throughout the thickness of the substrate, but may be concentrated at one or both surfaces, and may be substantially only present on one or both surfaces of the substrate.

5 Specific detackifiers include, but are not limited to, powders, such as talc powder, calcium carbonate, mica; starches, such as corn starch; lycopodium powder; mineral fillers, such as titanium dioxide; silica powder; alumina; metal oxides in general; baking powder; kieselguhr; and the like. Polymers and other additives having low surface energy may also be used,
10 including a wide variety of fluorinated polymers, silicone additives, polyolefins and thermoplastics, waxes, debonding agents known in the paper industry including compounds having alkyl side chains such as those having 16 or more carbons, and the like. Compounds used as release agents for molds and candle making may also be considered, as well as, dry lubricants
15 and fluorinated release agents.

 In one embodiment, the detackifier comprises polytetrafluorethylene (PTFE), such as PTFE telomer (KRYTOX[®] DF) compound, used in the PTFE release agent dry lubricant MS-122DF, marketed by Miller-Stephenson (Danbury, CT) as a spray product. For example, PTFE
20 particles may be applied by spray to one side of the substrate prior to winding of the pre-moistened wipes. In one embodiment, a detackifying agent is applied to only one surface of the substrate prior to winding into a roll.

 The wetting composition desirably contains less than about 25 weight percent of detackifying agents based on the total weight of the wetting composition. More desirably, the wetting composition contains from about
25 0.01 weight percent to about 10 weight percent of detackifying agents, more specifically about 5% or less. Even more specifically, the wetting composition contains from about 0.05 weight percent to about 2 weight percent of detackifying agents.

30 In addition to acting as a detackifying agent, starch compounds may also improve the strength properties of the pre-moistened wipes. For

example, it has been found that ungelled starch particles, such as hydrophilic tapioca starch, when present at a level of about 1% or higher by weight relative to the weight of the wetting composition, can permit the pre-moistened wipe to maintain the same strength at a lower salt concentration than is possible without the presence of starch. Thus, for example, a given strength can be achieved with 2% salt in the wetting composition in the presence of salt compared to a level of 4% salt being needed without starch. Starch may be applied by adding the starch to a suspension of laponite to improve the dispersion of the starch within the wetting composition.

Microparticulates

The wetting composition of the present invention may be further modified by the addition of solid particulates or microparticulates. Suitable particulates include, but are not limited to, mica, silica, alumina, calcium carbonate, kaolin, talc, and zeolites. The particulates may be treated with stearic acid or other additives to enhance the attraction or bridging of the particulates to the binder system, if desired. Also, two-component microparticulate systems, commonly used as retention aids in the papermaking industry, may also be used. Such two-component microparticulate systems generally comprise a colloidal particle phase, such as silica particles, and a water-soluble cationic polymer for bridging the particles to the fibers of the web to be formed. The presence of particulates in the wetting composition can serve one or more useful functions, such as (1) increasing the opacity of the pre-moistened wipes; (2) modifying the rheology or reducing the tackiness of the pre-moistened wipe; (3) improving the tactile properties of the wipe; or (4) delivering desired agents to the skin via a particulate carrier, such as a porous carrier or a microcapsule. Desirably, the wetting composition contains less than about 25 weight percent of particulate based on the total weight of the wetting composition. More specifically, the wetting composition may contain from about 0.05 weight percent to about 10 weight percent of microparticulate. Even more specifically, the wetting composition may

contain from about 0.1 weight percent to about 5 weight percent of microparticulate.

Microcapsules and Other Delivery Vehicles

5 Microcapsules and other delivery vehicles may also be used in the wetting composition of the present invention to provide skin-care agents; medications; comfort promoting agents, such as eucalyptus; perfumes; skin care agents; odor control additives; vitamins; powders; and other additives to the skin of the user. Specifically, the wetting composition may contain up to
10 about 25 weight percent of microcapsules or other delivery vehicles based on the total weight of the wetting composition. More specifically, the wetting composition may contain from about 0.05 weight percent to about 10 weight percent of microcapsules or other delivery vehicles. Even more specifically, the wetting composition may contain from about 0.2 weight percent to about
15 5.0 weight percent of microcapsules or other delivery vehicles.

 Microcapsules and other delivery vehicles are well known in the art. For example, POLY-PORE[®] E200 (Chemdal Corp., Arlington Heights, IL), is a delivery agent comprising soft, hollow spheres that can contain an additive at over 10 times the weight of the delivery vehicle.
20 Known additives reported to have been used with POLY-PORE[®] E200 include, but are not limited to, benzoyl peroxide, salicylic acid, retinol, retinyl palmitate, octyl methoxycinnamate, tocopherol, silicone compounds (DC 435), and mineral oil. Another useful delivery vehicle is a sponge-like material marketed as POLY-PORE[®] L200, which is reported to have been
25 used with silicone (DC 435) and mineral oil. Other known delivery systems include cyclodextrins and their derivatives, liposomes, polymeric sponges, and spray-dried starch.

 Additives present in microcapsules are isolated from the environment and the other agents in the wetting composition until the wipe is
30 applied to the skin, whereupon the microcapsules break and deliver their load to the skin or other surfaces.

Preservatives and Anti-Microbial Agents

The wetting composition of the present invention may also contain preservatives and/or anti-microbial agents. Several preservatives and/or anti-microbial agents, such as Mackstat H 66 (available from McIntyre Group, Chicago, IL), have been found to give excellent results in preventing bacteria and mold growth. Other suitable preservatives and anti-microbial agents include, but are not limited to DMDM hydantoin (*e.g.*, Glydant Plus™, Lonza, Inc., Fair Lawn, NJ), iodopropynyl butylcarbamate, Kathon (Rohm and Hass, Philadelphia, PA), methylparaben, propylparaben, 2-bromo-2-nitropropane-1,3-diol, benzoic acid, and the like. Desirably, the wetting composition contains less than about 2 weight percent on an active basis of preservatives and/or anti-microbial agents based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 1 weight percent of preservatives and/or anti-microbial agents. Even more desirably, the wetting composition contains from about 0.01 weight percent to about 0.5 weight percent of preservatives and/or anti-microbial agents.

Wetting Agents and Cleaning Agents

A variety of wetting agents and/or cleaning agents may be used in the wetting composition of the present invention. Suitable wetting agents and/or cleaning agents include, but are not limited to, detergents and nonionic, amphoteric, and anionic surfactants, especially amino acid-based surfactants. Amino acid-based surfactant systems, such as those derived from amino acids L-glutamic acid and other natural fatty acids, offer pH compatibility to human skin and good cleansing power, while being relatively safe and providing improved tactile and moisturization properties compared to other anionic surfactants. One function of the surfactant is to improve wetting of the dry substrate with the wetting composition. Another function of the surfactant can be to disperse bathroom soils when the pre-moistened wipe contacts a soiled

area and to enhance their absorption into the substrate. The surfactant can further assist in make-up removal, general personal cleansing, hard surface cleansing, odor control, and the like.

5 One commercial example of an amino-acid based surfactant is acylglutamate, marketed under the Amisoft name by Ajinomoto Corp., Tokyo, Japan. Desirably, the wetting composition contains less than about 3 weight percent of wetting agents and/or cleaning agents based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 2 weight percent of wetting agents
10 and/or cleaning agents. Even more desirably, the wetting composition contains from about 0.1 weight percent to about 0.5 weight percent of wetting agents and/or cleaning agents.

Although amino-acid based surfactants are particularly useful in the wetting compositions of the present invention, a wide variety of
15 surfactants may be used in the present invention. Suitable non-ionic surfactants include, but are not limited to, the condensation products of ethylene oxide with a hydrophobic (oleophilic) polyoxyalkylene base formed by the condensation of propylene oxide with propylene glycol. The hydrophobic portion of these compounds desirably has a molecular weight sufficiently high so as to render it water-insoluble. The addition of
20 polyoxyethylene moieties to this hydrophobic portion increases the water-solubility of the molecule as a whole, and the liquid character of the product is retained up to the point where the polyoxyethylene content is about 50% of the total weight of the condensation product. Examples of compounds of this type
25 include commercially-available Pluronic surfactants (BASF Wyandotte Corp.), especially those in which the polyoxypropylene ether has a molecular weight of about 1500-3000 and the polyoxyethylene content is about 35-55% of the molecule by weight, i.e. Pluronic L-62.

Other useful nonionic surfactants include, but are not limited
30 to, the condensation products of C_8 - C_{22} alkyl alcohols with 2-50 moles of ethylene oxide per mole of alcohol. Examples of compounds of this type

include the condensation products of C_{11} - C_{15} secondary alkyl alcohols with 3-50 moles of ethylene oxide per mole of alcohol, which are commercially-available as the Poly-Tergent SLF series from Olin Chemicals or the TERGITOL[®] series from Union Carbide, i.e. TERGITOL[®] 25-L-7, which is formed by condensing about 7 moles of ethylene oxide with a C_{12} - C_{15} alkanol.

Other nonionic surfactants, which may be employed in the wetting composition of the present invention, include the ethylene oxide esters of C_6 - C_{12} alkyl phenols such as (nonylphenoxy)polyoxyethylene ether. Particularly useful are the esters prepared by condensing about 8-12 moles of ethylene oxide with nonylphenol, i.e. the IGEPAL[®] CO series (GAF Corp.).

Further non-ionic surface active agents include, but are not limited to, alkyl polyglycosides (APG), derived as a condensation product of dextrose (D-glucose) and a straight or branched chain alcohol. The glycoside portion of the surfactant provides a hydrophile having high hydroxyl density, which enhances water solubility. Additionally, the inherent stability of the acetal linkage of the glycoside provides chemical stability in alkaline systems. Furthermore, unlike some non-ionic surface active agents, alkyl polyglycosides have no cloud point, allowing one to formulate without a hydrotrope, and these are very mild, as well as readily biodegradable non-ionic surfactants. This class of surfactants is available from Horizon Chemical under the trade names of APG-300, APG-350, APG-500, and APG-500.

Silicones are another class of wetting agents available in pure form, or as microemulsions, macroemulsions, and the like. One exemplary non-ionic surfactant group is the silicone-glycol copolymers. These surfactants are prepared by adding poly(lower)alkylenoxy chains to the free hydroxyl groups of dimethylpolysiloxanols and are available from the Dow Corning Corp as Dow Corning 190 and 193 surfactants (CTFA name: dimethicone copolyol). These surfactants function, with or without any volatile silicones used as solvents, to control foaming produced by the other surfactants, and also impart a shine to metallic, ceramic, and glass surfaces.

Anionic surfactants may also be used in the wetting compositions of the present invention. Anionic surfactants are useful due to their high detergency include anionic detergent salts having alkyl substituents of 8 to 22 carbon atoms such as the water-soluble higher fatty acid alkali metal soaps, e.g., sodium myristate and sodium palmitate. A preferred class of anionic surfactants encompasses the water-soluble sulfated and sulfonated anionic alkali metal and alkaline earth metal detergent salts containing a hydrophobic higher alkyl moiety (typically containing from about 8 to 22 carbon atoms) such as salts of higher alkyl mono or polynuclear aryl sulfonates having from about 1 to 16 carbon atoms in the alkyl group, with examples available as the Bio-Soft series, i.e. Bio-Soft D-40 (Stepan Chemical Co.).

Other useful classes of anionic surfactants include, but are not limited to, the alkali metal salts of alkyl naphthalene sulfonic acids (methyl naphthalene sodium sulfonate, Petro AA, Petrochemical Corporation); sulfated higher fatty acid monoglycerides such as the sodium salt of the sulfated monoglyceride of cocoa oil fatty acids and the potassium salt of the sulfated monoglyceride of tallow fatty acids; alkali metal salts of sulfated fatty alcohols containing from about 10 to 18 carbon atoms (e.g., sodium lauryl sulfate and sodium stearyl sulfate); sodium C₁₄-C₁₆-alphaolefin sulfonates such as the Bio-Terge series (Stepan Chemical Co.); alkali metal salts of sulfated ethyleneoxy fatty alcohols (the sodium or ammonium sulfates of the condensation products of about 3 moles of ethylene oxide with a C₁₂-C₁₅ n-alkanol, i.e., the Neodol ethoxysulfates, Shell Chemical Co.); alkali metal salts of higher fatty esters of low molecular weight alkylol sulfonic acids, e.g. fatty acid esters of the sodium salt of isothionic acid, the fatty ethanolamide sulfates; the fatty acid amides of amino alkyl sulfonic acids, e.g. lauric acid amide of taurine; as well as numerous other anionic organic surface active agents such as sodium xylene sulfonate, sodium naphthalene sulfonate, sodium toluene sulfonate and mixtures thereof.

A further useful class of anionic surfactants includes the 8-(4-n-alkyl-2-cyclohexenyl)-octanoic acids, wherein the cyclohexenyl ring is substituted with an additional carboxylic acid group. These compounds or their potassium salts, are commercially-available from Westvaco Corporation as Diacid 1550 or H-240. In general, these anionic surface active agents can be employed in the form of their alkali metal salts, ammonium or alkaline earth metal salts.

Macroemulsions and Microemulsion of Silicone Particles

The wetting composition may further comprise an aqueous microemulsion of silicone particles. For example, U.S. Pat. No. 6,037,407, "Process for the Preparation of Aqueous Emulsions of Silicone Oils and/or Gums and/or Resins" issued March 14, 2000, discloses organopolysiloxanes in an aqueous microemulsion. Desirably, the wetting composition contains less than about 5 weight percent of a microemulsion of silicone particles based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.02 weight percent to about 3 weight percent of a microemulsion of silicone particles. Even more desirably, the wetting composition contains from about 0.02 weight percent to about 0.5 weight percent of a microemulsion of silicone particles.

Silicone emulsions in general may be applied to the pre-moistened wipe by any known coating method. For example, the pre-moistened wipe may be moistened with an aqueous composition comprising a water-dispersible or water-miscible, silicone-based component that is compatible with the activating compound in the wetting composition. Further, the wipe can comprise a nonwoven web of fibers having a water-dispersible binder, wherein the web is moistened with a lotion comprising a silicone-based sulfosuccinate. The silicone-based sulfosuccinate provides gentle and effective cleansing without a high level of surfactant. Additionally, the silicone-based sulfosuccinate provides a solubilization function, which

prevents precipitation of oil-soluble components, such as fragrance components, vitamin extracts, plant extracts, and essential oils.

In one embodiment of the present invention, the wetting composition comprises a silicone copolyol sulfosuccinate, such as disodium dimethicone copolyol sulfosuccinate and diammonium dimethicone copolyolsulfosuccinate. Desirably, the wetting composition comprises less than about 2 percent by weight of the silicone-based sulfosuccinate, and more desirably from about 0.05 percent to about 0.30 percent by weight of the silicone-based sulfosuccinate.

In another example of a product comprising a silicone emulsions, Dow Corning 9506 powder may also be present in the wetting composition. Dow Corning 9506 powder is believed to comprise a dimethicone/vinyldimethicone cross-polymer and is a spherical powder, which is said to be useful in controlling skin oils (see "New Chemical Perspectives," *Soap and Cosmetics*, Vol. 76, No. 3, March 2000, p. 12). Thus, a water-dispersible wipe, which delivers a powder effective in controlling skin oil, is also within the scope of the present invention. Principles for preparing silicone emulsions are disclosed in WO 97/10100, published March 20, 1997.

Emollients

The wetting composition of the present invention may also contain one or more emollients. Suitable emollients include, but are not limited to, PEG 75 lanolin, methyl gluceth 20 benzoate, C₁₂-C₁₅ alkyl benzoate, ethoxylated cetyl stearyl alcohol, products marketed as Lambent wax WS - L, Lambent WD - F, Cetiol HE (Henkel Corp.), Glucam P20 (Amerchol), Polyox WSR N-10 (Union Carbide), Polyox WSR N-3000 (Union Carbide), Luviquat (BASF), Finsolv SLB 101 (Finetex Corp.), mink oil, allantoin, stearyl alcohol, Estol 1517 (Unichema), and Finsolv SLB 201 (Finetex Corp.).

An emollient can also be applied to a surface of the article prior to or after wetting with the wetting composition. Such an emollient may be

insoluble in the wetting composition and can be immobile except when exposed to a force. For example, a petrolatum-based emollient can be applied to one surface in a pattern, after which the other surface is wetted to saturate the wipe. Such a product could provide a cleaning surface and an opposing skin treatment surface.

The emollient composition in such products and other products of the present invention can comprise a plastic or fluid emollient such as one or more liquid hydrocarbons (e.g., petrolatum), mineral oil and the like, vegetable and animal fats (e.g., lanolin, phospholipids and their derivatives) and/or a silicone materials such as one or more alkyl substituted polysiloxane polymers, including the polysiloxane emollients disclosed in U.S. Pat. No. 5,891,126, issued Apr. 6, 1999 to Osborn, III et al. Optionally, a hydrophilic surfactant may be combined with a plastic emollient to improve wettability of the coated surface. In some embodiments of the present invention, it is contemplated that liquid hydrocarbon emollients and/or alkyl substituted polysiloxane polymers may be blended or combined with one or more fatty acid ester emollients derived from fatty acids or fatty alcohols.

In an embodiment of the present invention, the emollient material is in the form of an emollient blend. Desirably, the emollient blend comprises a combination of one or more liquid hydrocarbons (e.g., petrolatum), mineral oil and the like, vegetable and animal fats (e.g., lanolin, phospholipids and their derivatives), with a silicone material such as one or more alkyl substituted polysiloxane polymers. More desirably, the emollient blend comprises a combination of liquid hydrocarbons (e.g., petrolatum) with dimethicone or with dimethicone and other alkyl substituted polysiloxane polymers. In some embodiments of the present invention, it is contemplated that blends of liquid hydrocarbon emollients and/or alkyl substituted polysiloxane polymers may be blended with one or more fatty acid ester emollients derived from fatty acids or fatty alcohols. PEG-7 glyceryl cocoate, available as Standamul HE (Henkel Corp., Hoboken, N.J), can also be considered.

Water-soluble, self-emulsifying emollient oils, which are useful in the present wetting compositions, include the polyoxyalkoxylated lanolins and the polyoxyalkoxylated fatty alcohols, as disclosed in U.S. Pat. No. 4,690,821, issued Sep. 1, 1987 to Smith et al. The polyoxyalkoxy chains desirably will comprise mixed propylenoxy and ethyleneoxy units. The lanolin derivatives will typically comprise about 20-70 such lower-alkoxy units while the C₁₂ -C₂₀ - fatty alcohols will be derivatized with about 8-15 lower- alkyl units. One such useful lanolin derivative is Lanexol AWS (PPG-12-PEG-50, Croda, Inc., New York, N.Y.). A useful poly(15-20)C₂-C₃-alkoxylate is PPG-5-Ceteth-20, known as Procetyl AWS (Croda, Inc.).

According to one embodiment of the present invention, the emollient material reduces undesirable tactile attributes, if any, of the wetting composition. For example, emollient materials, including dimethicone, can reduce the level of tackiness that may be caused by the ion-sensitive binder or other components in the wetting composition, thus serving as a detackifier.

Desirably, the wetting composition contains less than about 25 weight percent of emollients based on the total weight of the wetting composition. More specifically, the wetting composition may comprise less than about 5 weight percent emollient, and most specifically less than about 2% emollient. More desirably, the wetting composition may contain from about 0.01 weight percent to about 8 weight percent of emollients. Even more desirably, the wetting composition may contain from about 0.2 weight percent to about 2 weight percent of emollients.

In one embodiment, the wetting composition and/or pre-moistened wipes of the present invention comprise an oil-in-water emulsion comprising an oil phase containing at least one emollient oil and at least one emollient wax stabilizer dispersed in an aqueous phase comprising at least one polyhydric alcohol emollient and at least one organic water-soluble detergent, as disclosed in U.S. Pat. No. 4,559,157, issued Dec. 17, 1985 to Smith et al., the entirety of which is herein incorporated by reference.

Surface Feel Modifiers

Surface feel modifiers are used to improve the tactile sensation (e.g., lubricity) of the skin during use of the product. Suitable surface feel modifiers include, but are not limited to, commercial debonders; and softeners, such as the softeners used in the art of tissue making including quaternary ammonium compounds with fatty acid side groups, silicones, waxes, and the like. Exemplary quaternary ammonium compounds with utility as softeners are disclosed in U.S. Pat. No. 3,554,862, issued to Hervey et al. on Jan. 12, 1971; U.S. Pat. No. 4,144,122, issued to Emanuelsson et al., Mar. 13, 1979, U.S. Pat. No. 5,573,637, issued to Ampulski et al. Nov. 12, 1996; and U.S. Pat. No. 4,476,323, issued to Hellsten et al., Oct. 9, 1984, the entirety of all of which is herein incorporated by reference. Desirably, the wetting composition contains less than about 2 weight percent of surface feel modifiers based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 1 weight percent of surface feel modifiers. Even more desirably, the wetting composition contains from about 0.01 weight percent to about 0.05 weight percent of surface feel modifiers.

Fragrances

A variety of fragrances may be used in the wetting composition of the present invention. Desirably, the wetting composition contains less than about 2 weight percent of fragrances based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 1 weight percent of fragrances. Even more desirably, the wetting composition contains from about 0.01 weight percent to about 0.05 weight percent of fragrances.

Fragrance Solubilizers

Further, a variety of fragrance solubilizers may be used in the wetting composition of the present invention. Suitable fragrance solubilizers

include, but are not limited to, polysorbate 20, propylene glycol, ethanol, isopropanol, diethylene glycol monoethyl ether, dipropylene glycol, diethyl phthalate, triethyl citrate, Ameroxol OE-2 (Amerchol Corp.), Brij 78 and Brij 98 (ICI Surfactants), Arlasolve 200 (ICI Surfactants), Calfax 16L-35 (Pilot Chemical Co.), Capmul POE-S (Abitec Corp.), Finsolv SUBSTANTIAL (Finetex), and the like. Desirably, the wetting composition contains less than about 2 weight percent of fragrance solubilizers based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 1 weight percent of fragrance solubilizers. Even more desirably, the wetting composition contains from about 0.01 weight percent to about 0.05 weight percent of fragrance solubilizers.

Opacifiers

Suitable opacifiers include, but are not limited to, titanium dioxide or other minerals or pigments, and synthetic opacifiers such as REACTOPAQUE[®] particles (available from Sequa Chemicals, Inc., Chester, South Carolina). Desirably, the wetting composition contains less than about 2 weight percent of opacifiers based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 1 weight percent of opacifiers. Even more desirably, the wetting composition contains from about 0.01 weight percent to about 0.05 weight percent of opacifiers.

pH Control Agents

Suitable pH control agents for use in the wetting composition of the present invention include, but are not limited to, malic acid, citric acid, hydrochloric acid, acetic acid, sodium hydroxide, potassium hydroxide, and the like. An appropriate pH range minimizes the amount of skin irritation resulting from the wetting composition on the skin. Desirably, the pH range of the wetting composition is from about 3.5 to about 6.5. More desirably, the

pH range of the wetting composition is from about 4 to about 6. Desirably, the wetting composition contains less than about 2 weight percent of a pH adjuster based on the total weight of the wetting composition. More desirably, the wetting composition contains from about 0.01 weight percent to about 1 weight percent of a pH adjuster. Even more desirably, the wetting composition contains from about 0.01 weight percent to about 0.05 weight percent of a pH adjuster.

Although a variety of wetting compositions, formed from one or more of the above-described components, may be used with the wet wipes of the present invention, in one embodiment, the wetting composition contains the following components, given in weight percent of the wetting composition, as shown in Table 1 below:

Table 1. Wetting Composition Components

<i>Wetting Composition Component:</i>	<i>Weight Percent:</i>
Deionized Water	about 86 to about 98
Activating compound	about 2 to about 20
Preservative	Up to about 2
Surfactant	Up to about 2
Silicone Emulsion	Up to about 1
Emollient	Up to about 1
Fragrance	Up to about 0.3
Fragrance solubilizer	Up to about 0.5
pH adjuster	Up to about 0.2

In another embodiment of the present invention, the wetting composition comprises the following components, given in weight percent of the wetting composition, as shown in Table 2 below:

Table 2 -- Wetting Composition Components

<i>Class of Wetting Composition Component:</i>	<i>Specific Wetting Composition Component:</i>	<i>Component Name:</i>	<i>Weight Percent:</i>
Vehicle	Deionized Water		about 86 to about 98
Activating compound	Sodium Chloride (Millport Ent., Milwaukee, WI)		about 2 to about 20
Preservative	Glycerin, IPBC and DMDM Hydantoin	Mackstat H-66 (McIntyre Group, Chicago, IL)	Up to about 2
Surfactant	Acyl Glutamate	CS22 (Ajinomoto, Tokyo, Japan)	Up to about 2
Silicone Emulsion (Detackifier/Skin Feel agent)	Dimethiconol and TEA Dodecylbenzene Sulfonate	DC1785 (Dow Corning, Midland, MI)	Up to about 1
Emollient	PEG-75 Lanolin	Solulan L- 575 (Amerchol, Middlesex, NJ)	Up to about 1
Fragrance	Fragrance	Dragoco 0/708768 (Dragoco, Roseville, MN)	Up to about 0.3
Fragrance solubilizer	Polysorbate 20	Glennsurf L20 (Glenn Corp., St. Paul, MN)	Up to about 0.5
pH adjuster	Malic Acid to pH 5 (Haarman & Reimer, Tetrboro, NJ)		Up to about 0.2

In another embodiment of the present invention, the wetting composition comprises the following components, given in weight percent of the wetting composition, as shown in Table 3 below:

Table 3 -- An Exemplary Wetting Composition

<i>Class of Wetting composition</i> <i>Component:</i>	<i>Specific Wetting composition</i> <i>Component:</i>	<i>Component Name:</i>	<i>Weight Percent:</i>
Vehicle	Deionized Water		about 93
Activating compound	Sodium Chloride		about 4
Preservative	Glycerin, IPBC and DMDM Hydantoin	Mackstat H-66	about 1
Surfactant	Acyl Glutamate	CS22/ECS 22P	about 1
Silicone Emulsion	Dimethiconol and TEA Dodecylbenzene Sulfonate	DC 1784/ DC1785	about 0.5
Emollient	PEG-75 Lanolin	Solulan L- 575	about 0.25
Fragrance	Fragrance	Dragoco Fragrance 0/708768	about 0.05
Fragrance solubilizer	Polysorbate 20	Glennsurf L20	about 0.25
pH adjuster	Malic Acid to pH 5		about 0.07

It should be noted that the above-described wetting compositions of the present invention may be used with any one of the above-described ion-sensitive binder compositions of the present invention. Further, the above-described wetting compositions of the present invention may be used with any other binder composition, including conventional binder compositions, or with any known fibrous or absorbent substrate, whether dispersible or not.

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Strength Properties

Unless otherwise specified, tensile testing is performed according to the following protocol. Testing of dry product should be conducted under Tappi conditions (50% relative humidity, 73°F) with a procedure similar to ASTM-1117-80, section 7. Tensile tests are performed with a constant crosshead speed tensile tester such as the Thwing Albert 1256-100 tensile tester with an RSA-2 10-kg load cell. Specimens are cut to 3-inch widths and 6 inch lengths, and mounted between jaws with a 4-inch gauge length. The crosshead speed is 12 inches per minute. Peak load (for tensile strength) and elongation at peak load (for stretch) are measured. For cross direction (CD) tensile tests, the sample is cut in the cross direction. For machine direction (MD) tensile tests, the sample is cut in the cross direction.

Tensile tests in the dry state are reported for webs taken prior to application of the wetting composition. The machine direction dry tensile strength is abbreviated as "MDDT," and the cross direction dry tensile strength as "CDDT." The results can be reported as kg/3-in or converted to units of g/in or g/2.54 cm.

Based on the dry weight of the specimen cut to the appropriate size, an excess amount of wetting solution (4% saline solution with no other additives, unless otherwise specified) is applied to reach a solution add-on of 250-400%. The wetted specimens are then immediately passed through an Atlas Lab Wringer (Atlas Electric Devices Company, Chicago, Ill. No. 10404 LW-1, no load) to uniformly distribute the solution in the sample and gently remove the excess solution to achieve a final solution add-on of 200%. Several iterations or passes may be needed to reach the add-on target depending on the sample. The completed, pre-moistened samples are then bagged in plastic to prevent dry-out before testing.

Cross direction wet tensile tests (CDWT) or machine direction wet tensile strength (MDWT) are performed as described above using the pre-

moistened sample as is, after the sample has equilibrated by sitting overnight in a sealed plastic bag.

For tests related to strength loss in a pre-moistened web occurring after exposure to a new solution, a container having dimensions of 200 mm by 120 mm and deep enough to hold 1000 ml is filled with 700 ml of the selected soak solution. No more than 108 square inches of sample are soaked in the 700 ml of soaking solution, depending on specimen size. The pre-moistened specimens, that have equilibrated overnight, are immersed in the soak solution and then allowed to soak undisturbed for a specified time period (typically 1 hour). At the completion of the soak period, samples are carefully retrieved from the soak solution, allowed to drain, and then tested immediately as described above (*i.e.*, the sample is immediately mounted in the tensile tester and tested, without being passed through the wringer). In cases with highly dispersible materials, the samples often cannot be retrieved from the soaking solution without falling apart. The soaked tensile values for such samples are recorded as zero for the corresponding solution.

For the deionized soaked cross-direction wet tensile test, S-CDWT, the sample is immersed in deionized water for 1 hour and then tested. For the hard-water soaked cross-direction wet tensile test, S-CDWT-M (M indicating divalent metal ions), the sample is immersed in water containing 200 ppm of $\text{Ca}^{++}/\text{Mg}^{++}$ in a 2:1 ratio prepared from calcium chloride and magnesium chloride, soaked for one hour and then tested. For the medium hard water soaked cross-direction wet tensile test, MS-CDWT-M, the sample is immersed in water containing 50 ppm of $\text{Ca}^{++}/\text{Mg}^{++}$ in a 2:1 ratio, soaked for one hour and then tested. Testing done with other time increments or soaking solutions should be so indicated to prevent confusion with the S-CDWT or S-CDWT-M tests.

In one embodiment of the present invention, wet wipes are produced using the above-described wetting composition in Table 3 and an air-laid fibrous material comprising about 80 weight percent of bleached kraft fibers and 20 weight percent of any of the above-described ion-sensitive

binder compositions of the present invention, wherein the weight percentages are based on the total weight of the dry nonwoven fabric. In a further embodiment of the present invention, wet wipes are produced using the above-described wetting composition in Table 3 and an air-laid fibrous material comprising 90 weight percent of softwood fibers and 10 weight percent of an ion-sensitive binder of the present invention. The amount of wetting composition added to the nonwoven fabric, relative to the weight of the dry nonwoven fabric in these embodiments, is desirably about 180 percent to about 240 weight percent.

Desirably, the wet wipes of the present invention possess an in-use wet tensile strength (CDWT) of at least 100 g/in, and a tensile strength of less than about 30 g/in after being soaked in water having a concentration of Ca^{2+} and/or Mg^{2+} ions of about 50 ppm for about one hour (MS-CDWT-M). More desirably, the wet wipes possess an in-use wet tensile strength of at least 300 g/in (CDWT), and a tensile strength of less than about 30 g/in after being soaked in water having a concentration of Ca^{2+} and/or Mg^{2+} ions of about 50 ppm for about one hour (MS-CDWT-M). In a further embodiment, the wet wipes desirably possess an in-use wet tensile strength of at least 200 g/in (CDWT), and a tensile strength of less than about 20 g/in after being soaked in water having a concentration of Ca^{2+} and/or Mg^{2+} ions of about 200 ppm for about one hour (S-CDWT-M). Even more desirably, the wet wipes possess an in-use wet tensile strength of at least 300 g/in, and a tensile strength of less than about 20 g/in after being soaked in water having a concentration of Ca^{2+} and/or Mg^{2+} ions of about 200 ppm for about one hour (S-CDWT-M).

Desirably, the wet wipes treated with the binder material of the present invention possess an in-use wet tensile strength of at least 100 g/in for a 1 inch width sample in the cross machine direction when soaked with 10% to 400% by weight wet wipes solution containing more than 2% by weight monovalent ion (NaCl) concentration and a tensile strength of less than about 30 g/in after being soaked in deionized water for about one hour. More desirably, the wet wipes treated with the binder material of the present

invention possess an in-use tensile strength of at least 200 g/in for a 1 inch width sample in the cross machine direction when soaked with 10% to 400% by weight wet wipes solution containing more than 2% by weight monovalent ion (NaCl) concentration and a tensile strength of less than about 30 g/in after being soaked in deionized water for about one hour. More desirably, the wet wipes treated with the binder material of the present invention possess an in-use tensile strength of at least 300 g/in for a 1 inch width sample in the cross machine direction when soaked with 10% to 400% by weight wet wipes solution containing more than 2% by weight monovalent ion (NaCl) concentration and a tensile strength of less than about 20 g/in after being soaked in deionized water for about one hour.

Products with high basis weights or wet strengths than flushable wet wipes may have relatively higher wet tensile strength. For example, products, such as pre-moistened towels or hard-surface cleaning wipes, may have basis weights above 70 gsm, such as from 80 gsm to 150 gsm. Such products can have CDWT values of 500 g/in or greater, with S-CDWT values of about 150 g/in or less, more specifically about 100 g/in or less, and most specifically about 50 g/in or less, with similar ranges possible for S-CDWT-M.

Dispersibility

Prior efforts to measure dispersibility of webs, whether dry or pre-moistened, have commonly relied on systems in which the web was exposed to shear while in water, such as measuring the time for a web to break up while being agitated by a mechanical mixer. The constant exposure to shear offers an unrealistic and overly optimistic test for products designed to be flushed in a toilet, where the level of shear is weak and extremely brief. Once the product has passed through the neck of the toilet and entered a septic tank, shear rates may be negligible. Further, the product may not be fully wetted with water from the toilet bowl when it is flushed, or rather, there may not have been adequate time for the wetting composition of the product to

have been replaced with the water of the toilet bowl when the momentary shear of flushing is applied. Thus, previous measurements of dispersibility could suggest that a product is dispersible when, in fact, it may be poorly suited for septic system.

5 For a realistic appraisal of dispersibility, it is believed that a relatively static measure is needed to better simulate the low shear that real products will experience once they have become fully wetted with water from the toilet. Thus, a test method for dispersibility has been developed which does not rely on shear and which provides an improved means of assessing
10 suitability of a product for a septic system. In this method, the tensile strength of a product is measured in its original, wetted form (the CDWT measurement described above) and after the product has been soaked in a second solution for one hour (either the S-CDWT or S-CDWT-M test). The second solution can be either deionized water for determination of the "Deionized
15 Dispersibility" value or hard water (according to the S-CDWT-M test) for determination of the "Hard Water Dispersibility" value. In either case, the Dispersibility is defined as (1 minus the ratio of the cross-direction wet tensile strength in the second solution divided by the original cross-direction wet tensile strength) * 100%. Thus, if a pre-moistened wipe loses 75% of its CD
20 wet tensile strength after soaking in hard water for one hour, the Hard Water Dispersibility is $(1-0.25)*100\% = 75\%$. The articles of the present invention can have a Deionized Dispersibility of 80% or greater, more specifically 90% or greater, specifically still 95% or greater, and can have a Deionized Dispersibility of about 100%. The articles of the present invention can have a
25 Hard Water Dispersibility of 70% or greater, more specifically 80% or greater, specifically still about 90% or greater, and can have a Deionized Dispersibility of about 100%.

Method of Making Wet Wipes

30 The pre-moistened wipes of the present invention can be made in several ways. In one embodiment, the ion-sensitive polymer composition is

applied to a fibrous substrate as part of an aqueous solution or suspension, wherein subsequent drying is needed to remove the water and promote binding of the fibers. In particular, during drying, the binder migrates to the crossover points of the fibers and becomes activated as a binder in those regions, thus providing acceptable strength to the substrate. For example, the following steps can be applied:

1. Providing an absorbent substrate that is not highly bonded (e.g., an unbonded airlaid, a tissue web, a carded web, fluff pulp, etc.).

2. Applying an ion-sensitive polymer composition to the substrate, typically in the form of a liquid, suspension, or foam.

3. Drying the substrate to promote bonding of the substrate. The substrate may be dried such that the peak substrate temperature does not exceed about 100° to 160° C. In one embodiment, the substrate temperature does not exceed 60° C to 80° C.

5. Applying a wetting composition to the substrate.

6. Placing the wetted substrate in roll form or in a stack and packaging the product.

Application of the ion-sensitive polymer composition to the substrate can be by means of spray; by foam application; by immersion in a bath; by curtain coating; by coating and metering with a wire-wound rod; by passage of the substrate through a flooded nip; by contact with a pre-metered wetted roll coated with the binder solution; by pressing the substrate against a deformable carrier containing the ion-sensitive polymer composition such as a sponge or felt to effect transfer into the substrate; by printing such as gravure, inkjet, or flexographic printing; and any other means known in the art.

In the use of foams to apply a binder or co-binder polymer, the mixture is frothed, typically with a foaming agent, and spread uniformly on the substrate, after which vacuum is applied to pull the froth through the substrate. Any known foam application method can be used, including that of U.S. Pat. No. 4,018,647, "Process for the Impregnation of a Wet Fiber Web with a Heat Sensitized Foamed Latex Binder," issued Apr. 19, 1977 to

Wietsma, the entirety of which is herein incorporated by reference. Wietsma discloses a method wherein a foamed latex is heat-sensitized by the addition of a heat-sensitizer such as functional siloxane compounds including siloxane oxyalkylene block copolymers and organopolysiloxanes. Specific examples of applicable heat-sensitizers and their use thereof for the heat sensitization of latices are described in the U.S. Pat. Nos. 3,255,140; 3,255,141; 3,483,240 and 3,484,394, all of which are incorporated herein by reference. The use of a heat-sensitizer is said to result in a product having a very soft and textile-like hand compared to prior methods of applying foamed latex binders.

The amount of heat-sensitizer to be added is dependent on, inter alia, the type of latex used, the desired coagulation temperature, the machine speed and the temperatures in the drying section of the machine, and will generally be in the range of about 0.05 to about 3% by weight, calculated as dry matter on the dry weight of the latex; but also larger or smaller amounts may be used. The heat sensitizer can be added in such an amount that the latex will coagulate far below the boiling point of water, for instance at a temperature in the range of 35° C. to 95° C., or from about 35° C. to 65° C.

Without wishing to be bound by theory, it is believed that a drying step after application of the binder solution and before application of the wetting composition enhances bonding of a fibrous substrate by driving the binder to fiber crossover points as moisture is driven off, thus promoting efficient use of the binder. However, in an alternative method, the drying step listed above is skipped, and the ion-sensitive polymer composition is applied to the substrate followed by application of the wetting composition without significant intermediate drying. In one version of this method, the ion-sensitive polymer composition selectively adheres to the fibers, permitting excess water to be removed in an optional pressing step without a significant loss of the binder from the substrate. In another version, no significant water removal occurs prior to application of the wetting composition. In yet another alternative method, the ion-sensitive polymer composition and the wetting

composition are applied simultaneously, optionally with subsequent addition of salt or other activating compounds to activate or further activate the binder.

The present invention is further illustrated by the following examples, which are not to be construed in any way as imposing limitations upon the scope thereof. On the contrary, it is to be clearly understood that resort may be had to various other embodiments, modifications, and equivalents thereof which, after reading the description herein, may suggest themselves to those skilled in the art without departing from the spirit of the present invention and/or the scope of the appended claims.

As used herein, the "thickness" of a web is measured with a 3-in acrylic plastic disk connected to the spindle of a Mitutoyo Digimatic Indicator (Mitutoyo Corporation, 31-19, Shiba 5-chome, Minato-ku, Tokyo 108, Japan) and which delivers a net load of 0.05 psi to the sample being measured. The Mitutoyo Digimatic Indicator is zeroed when the disk rests on a flat surface. When a sample having a size at least as great as the acrylic disk is placed under the disk, a thickness reading can be obtained from the digital readout of the indicator. Water-dispersible substrates of the present invention can have any suitable thickness, such as from about 0.1 mm to 5 mm. For wet wipes, thicknesses can be in the range of 0.2 mm to about 1 mm, more specifically from about 0.3 mm to about 0.7 mm. Thickness can be controlled, for example, by the application of compaction rolls during or after web formation, by pressing after binder or wetting composition has been applied, or by controlling the tension of winding when forming a roll good.

The use of the platen method to measure thickness gives an average thickness at the macroscopic level. Local thickness may vary, especially if the product has been embossed or has otherwise been given a three-dimensional texture.

EXAMPLE 1

Polymers were synthesized by free radical polymerization of varying combinations of the following monomers: acrylic acid, acrylamide,

butyl acrylate, 2-ethylhexyl acrylate and [2(methacryloyloxy)ethyl] trimethyl ammonium chloride ("MEMAC"). Each polymerization was conducted in methanol. A typical procedure is stated below.

Acrylamide (39.1 g, 0.55 mol), butyl acrylate (32.0 g, 0.25 mol), 2-ethylhexyl acrylate (18.4 g, 0.10 mol), and MEMAC (27.6 g of 75 wt% solution, 0.10 mol) were dissolved in 50 g of methanol. A free radical initiator, 2,2'-Azobisisobutyronitrile ("AIBN")² (0.66 g, 4.0 x 10⁻³ mol) was dissolved in 20 ml of methanol. The monomer solution was deoxygenated by bubbling N₂ through the solution for 20 minutes. To a 1000 ml round bottom, three neck flask equipped with a condenser, two addition funnels and a magnetic stirrer was added 125 g of methanol. The solvent was heated to gentle reflux under nitrogen. Monomers and initiator were added simultaneously from the addition funnels over a period of two hours. Polymerization was allowed to proceed for an additional two hours, at the end of which the addition funnels and condenser were replaced with a distillation head and a mechanical stir rod to remove methanol. A steady stream of N₂ was kept during distillation. When the distillation was completed (about 3 hours), 400 g of deionized water was added to the polymer solution. The heat was removed and the solution was allowed to stir overnight.

Alternatively, the polymers can be made by adding the monomers and initiator to the reaction flask all at once and reacting for four hours. This synthesis method is referred as "one pot" synthesis in the subsequent section. A total of eight polymers were synthesized and their compositions are summarized in Table 4 below.

Table 4 -- Polymer Composition

Sample	%MEMAC	%AM	%AA	%BA	%EHA
1	10	0	55	25	10
2	10	55	0	25	10
3	60	0	0	20	20
4	45	0	0	35	20
5	35	0	0	35	30
6	30	0	0	35	35

² What is the commercial source and brand name of this compound?

7	20	0	0	40	40
8	15	0	0	45	40

Sample preparation:

A water-dispersible, wet-laid nonwoven composed of BFF rayon fibers (1.5 d x 25 mm) was utilized as the base sheet for testing. Each base sheet was cut to an approximate size of 5.5 in (CD) x 9 in (MD). A piece of release paper was placed onto a notepad, followed by a base sheet. Both pieces were taped to the notepad with a single piece of Scotch tape. A #20 grooved, wire-wound rod was laid across the top of the sample. A strip of the polymer solution to be tested was poured along the rod. The rod was then rolled down the length of the sample, with gentle pressure applied. Excess polymer was wiped off the bottom of the release paper, and the sample was placed into a forced air oven at 600° C. for at least 10 minutes. The rod was cleaned between each sample as necessary. Once the samples were dry, they were removed from the oven. The top part of each sample was removed with a paper cutter. Each sample was then peeled from the release paper and the excess polymer film was gently pulled from the edges of the sample. Each sample sheet was then cut into ten 1 in (CID) x 4 in (MID) strips.

Tensile testing:

The SinTech 1/D tensile tester with Testworks 3.03 version software was utilized for all sample testing. All testing was conducted in the machine direction using a 50 pound load cell and pneumatic, rubberized grips. The gage length was set at 3 in, and the crosshead speed was 12 in/min. The wet samples were secured in the grips and stretched to failure. The peak load of each sample was recorded as the data of interest. The data was not normalized to a 100% add-on level. A value of "0" was entered for the peak load if the sample was determined to be dispersed. Samples were considered dispersed if individual strips could not be removed from the salt solution intact due to lack of structural integrity.

The in-use strength of each sample was simulated by soaking the tensile samples in various salt solutions. The concentrations of the salt solutions were chosen based upon a 2 wt% NaCl solution, which is equivalent to 0.34 M. The 0.68 M and 1.36 M NaCl solutions correspond to 4 wt% and 8 wt%, respectively. The same molarities were chosen for each of the other salts tested, though the weight percents are not necessarily equivalent to the sodium chloride solutions. Salt solutions tested included 0.34 M NaCl, 0.68 M NaCl, 1.36 M NaCl, 0.34 M CaCl₂, 0.68 M CaCl₂, 1.36 M CaCl₂, 0.34 M Na₂SO₄, 0.68 M Na₂SO₄, 1.36 M Na₂SO₄, 0.17 M ZnCl₂, 0.34 M ZnCl₂, 0.51 M ZnCl₂, 0.34 M ZnSO₄, 0.17 M ZnCl₂ + 0.17 M NaCl, 0.34 M LiCl, 0.34 M KCl, 0.34 M Na₃PO₄, and 0.34 MMgCl₂.

Twenty-four tensile samples were placed into the salt solution to be tested and allowed to soak overnight. Strips were added to the salt solution one at a time in order to avoid sticking the samples together. Average soaking time was approximately 17 hours, and volume of soaking solution was held constant at approximately 500 mL. Following the overnight soak, eight samples were tested directly to determine the peak load. This test simulated storage and in-use strength. Eight samples were placed into 200 ppm Ca²⁺/Mg²⁺ for 1 hour, and eight samples were placed into 200 ppm Ca²⁺/Mg²⁺ for 3 hours. The peak load of the samples was measured following the soaking times. This test simulated disposal in the hardest water found in the United States.

Trigger property:

Polymer 1 showed salt sensitivity and significant strength in several solutions of sodium chloride. However, the polymer failed to disperse in 200 ppm Ca²⁺/Mg²⁺, as shown in Table 5 below.

Table 5 -- Tensile strength of acrylic acid based polymer (g/in)

Salt Concentration	Overnight soak	1 hr. in hard water	3 hr. in hard water
0.34 M	565	606	648
0.68 M	639	649	634

1.36 M	722	600	608
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It is suspected that the positive charge on the polymer promotes dissociation of carboxylic acid groups creating carboxylate anions. Complex formation between the carboxylate anion and Ca^{2+} leads to cross-linking, which prevents dispersion in hard water. In order to obtain hard water dispersibility, the acrylic acid was replaced with acrylamide, a water-soluble monomer. The acrylamide polymer (sample 2) showed trigger behavior in several salt solutions as summarized in Table 6 below.

Table 6 -- Tensile strength of acrylamide based polymer in various salt solutions (g/in)

Salt Solution	Overnight Soak	1 hr. in Hard Water	3 hr. in Hard Water
0.34M NaCl	311	0	0
0.68M NaCl	317	0	0
1.36M NaCl	429	0	0
0.34M CaCl_2	664	636	656
0.68M CaCl_2	635	662	666
1.36M CaCl_2	603	616	596
0.34M Na_2SO_4	289	0	0
0.68M Na_2SO_4	486	160	0
1.36M Na_2SO_4	897	265	181
0.17M ZnCl_2	345	0	0
0.34M ZnCl_2	519	0	0
1.36M ZnCl_2	661	0	0

General trends showed that increasing the concentration of the salt solution increased the in-use strength of the polymer binder. The performance of the polymer is salt-specific. The polymer showed moderate strength and good dispersibility in NaCl. The strength of the polymer increased almost linearly with increasing Na_2SO_4 concentration, but reached good strength only at higher salt level. The dispersibility went in the opposite direction. The polymer showed nice strength in ZnCl_2 and good dispersibility. For CaCl_2 , the polymer showed good strength, independent of salt concentration in the range of investigation, but no strength loss was observed when placed in hard water. Both results suggested the cross-linking of the polymer caused by Ca^{2+} . Acrylamide has been generally regarded as salt insensitive, and its interaction

with Ca^{2+} is unknown. Based on the fact that the polymer dispersed well in hard water when soaked in other salts, such as ZnCl_2 and NaCl , it is clear that retardation only occurs above certain Ca^{2+} concentrations.

One concern with acrylamide based polymer is the toxicity of the residual acrylamide monomer in the polymer solution. To avoid the safety concern, acrylamide was removed from the composition. Sample 3--8 were made with only three monomers: MEMAC, BA and EHA. Table 7 summarized their trigger behavior.

Table 7 -- Trigger property of polymers of MEMAC, BA and EHA

Sample	3	4	5	6	7	8
Solubility	soluble	soluble	dispersed	dispersed	triggered	insoluble

They differ greatly in the salt sensitivity as small changes in composition can have large effects on the behavior of the polymer in solution. In general, their solubility decreases with decreasing MEMAC content. Samples 3 and 4 were soluble in water even in very high salt concentrations. Samples 5 and 6 were soluble in water and precipitated at high salt level, but showed no strength. Sample 8 is insoluble in water due to high hydrophobicity. Sample 7 had the best trigger property for its delicate balance between the hydrophobic and hydrophilic monomers. The behavior of the polymer in various salt solutions is shown in Table 8 below.

Table 8 -- Performance of polymer 7 in various salt solutions (g/in)

Salt Solution	Overnight Soak in Salt Solution	1 hr. Soak in Hard Water	3 hr. Soak in Hard Water
0.34M LiCl	65	0	0
0.34M NaCl	66	0	0
0.34M KCl	81	0	0
0.34M MgCl ₂	96	0	0
0.34M CaCl ₂	183	0	0
0.34M ZnCl ₂	861	33	25
0.17M ZnCl ₂ + 0.17M NaCl	819	27	12
0.065M ZnCl ₂ + 0.15M NaCl	0370	0	0
0.34M Na ₂ SO ₄	0	0	0
0.34M Na ₃ PO ₄	895	736	699

The polymer displayed trigger property in monovalent salts, namely LiCl, NaCl and KCl, but it lacked any significant strength, falling far short of the minimum goal of 300 g/in for in-use strength. The performance in divalent cations was highly selective. The polymer showed unremarkable strength in MgCl₂ and CaCl₂, but displayed excellent strength of over 800 g/in in ZnCl₂. The polymer also had excellent triggerability in mixed salt of 0.17M ZnCl₂ and 0.17 M NaCl. When the ZnCl₂ concentration was reduced to 0.065M, the strength was reduced. The polymer showed good hard water dispersibility from all those salts.

The effect of anions, especially multivalent anions, on the trigger property was tested. The strength of the polymer in NaCl is unremarkable, though the dispersibility is good. The polymer possessed zero strength in Na₂SO₄. Following this discovery, the polymer was also tested in ZnSO₄, in order to compare the exceptional strength of the polymer in Zn²⁺ ion to its complete lack of strength in SO₄²⁻ ions. The polymer displayed zero strength in ZnSO₄ solution. This surprising result is at once both frustrating and encouraging. Sulfate ion is known to be a good salt for insolubilizing anionic triggered polymers. However, it cannot be used for cationic triggers. On the positive side, however, the sulfate ion, which is common in waste water streams, will not interfere with the dispersion of the polymer. The polymer forms a non-dispersible complex with phosphate ions, as it would not even

disperse after several hours in deionized water. Fortunately, phosphate ion is extremely rare in waste water streams, and thus should pose no problem for the dispersibility of the polymer once in a product application.

5 The "one pot" polymer, which contains 20% MEMAC, 40% BA and 40% EHA, displayed excellent strength of 896.6 g/in following the overnight soak. However, the dispersibility of the "one pot" polymer was not as good as that of the model polymer, as it still displayed 138.7 g/in of strength after one hour in hard water, and 90.2 g/in of strength after a three hour soak. Overall, the "one pot" polymer displays good trigger property.

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EXAMPLE 2³

15 It should be understood, of course, that the foregoing relates only to certain disclosed embodiments of the present invention and that numerous modifications or alterations may be made therein without departing from the spirit and scope of the invention as set forth in the appended claims.

³ Need more test data.